

**INFORMATION
DISCLOSURE
STATEMENT**

Atty. Docket No.: 249.00020101

Serial No.: 09/970,318

Applicant(s): Cosgrove

Confirmation No.: 1885

Filing Date: October 3, 2001

Group: 1641

U.S. PATENT DOCUMENTS

Document Number	Date	Name	Class	Subclass	Filing Date If Appropriate
none					

FOREIGN PATENT DOCUMENTS

Examiner Initial	Document Number	Date	Country	Class	Subclass	Translation	
						Yes	No
	none						

OTHER DOCUMENTS (Including Authors, Title, Date, Pertinent Papers, etc.)

Examiner Initial	Document Description
MY	Adato et al., "Three novel mutations and twelve polymorphisms identified in the USH2A gene in Israeli USH2 families," <i>Hum Mutat. (Mutation in Brief)</i> ; published online for subscribers 2000 Mar 22 (6 pgs.).
MY	Adato et al., "Three novel mutations and twelve polymorphisms identified in the USH2A gene in Israeli USH2 families," <i>Hum Mutat.</i> 2000 Apr;15(4):abst. 388.
MY	Barkalow et al., "Localization of the major heparin-binding site in fibronectin," <i>J Biol Chem.</i> 1991 Apr 25;266(12):7812-8.
MY	Beck et al., "Structure and function of laminin: anatomy of a multidomain glycoprotein," <i>FASEB J.</i> 1990 Feb 1;4(2):148-60.
MY	Bork et al., "Structure and distribution of modules in extracellular proteins," <i>Q Rev Biophys.</i> 1996 May;29(2):119-67.
MY	Boughman et al., "Usher syndrome: definition and estimate of prevalence from two high-risk populations," <i>J Chronic Dis.</i> 1983;36(8):595-603.
MY	Bowditch et al., "Identification of a novel integrin binding site in fibronectin. Differential utilization by $\beta 3$ integrins," <i>J Biol Chem.</i> 1994 Apr 8;269(14):10856-63.

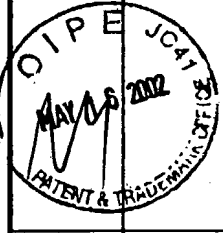
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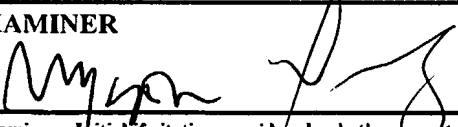
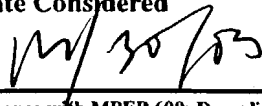
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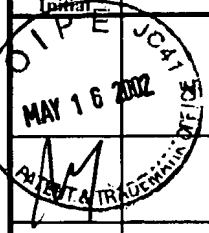
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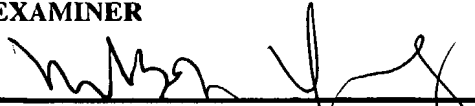
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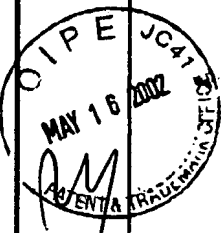
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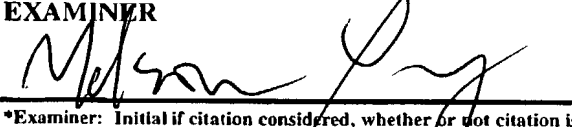
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M	National Center for Biotechnology Information, National Library of Medicine, National Institutes of Health, GenBank Locus HSUSH2A01, Accession No. AF091873, "Homo sapiens Usher syndrome type IIa protein gene, exons 1 and 2," [online]. Bethesda, MD [retrieved on 2002-04-05]. Retrieved from the Internet: <URL:http://www.ncbi.nlm.nih.gov/entrez/viewer.fcgi?query_key=4&db=nucleotide&page=0&dispmax=20&WebEnv=Wg%3Cn_FDG%5DE%60%3E%3D%3C%5DPGDJc_gTB%5EjbFkl%3C_JEH%3Dzcc%3EF%5EFJdTI%3D%3F%3C&WebEnvRq=1>; 3 pgs.

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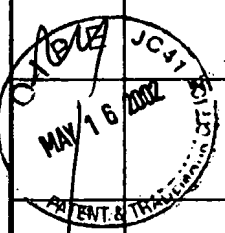
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MAY 20 2002

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	National Center for Biotechnology Information, National Library of Medicine, National Institutes of Health, GenBank Locus HSUSH2A17, Accession No. AF091889, "Homo sapiens Usher syndrome type IIa protein gene, exon 21 and complete cds," [online]. Bethesda, MD [retrieved on 2002-04-05]. Retrieved from the Internet: <URL: http://www.ncbi.nlm.nih.gov/entrez/viewer.fcgi?query_key=6&db=nucleotide&page=0&dispmax=20&WebEnv=C%5B%60%3CIgCAA%5C%5Ee%3D%3FHCK%3CCDB%3CdIEjhCdbaiH%3F%40AARqjH%5EGe%5E%3EkDFH&WebEnvRq=1 >; 3 pgs.
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	van der Loop et al., "Identification of COL4A5 defects in Alport's syndrome by immunohistochemistry of skin," <i>Kidney Int.</i> 1999 Apr;55(4):1217-24.
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